

NMR Backbone Dynamics of VEK-30 Bound to the Human Plasminogen Kringle 2 Domain

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ABSTRACT To gain insights into the mechanisms for the tight and highly specific interaction of the kringle 2 domain of human plasminogen (K2_{Pg}) with a 30-residue internal peptide (VEK-30) from a group A streptococcal M-like protein, the dynamic properties of free and bound K2_{Pg} and VEK-30 were investigated using backbone amide ¹⁵N-NMR relaxation measurements. Dynamic parameters, namely the generalized order parameter, S^2 , the local correlation time, τ_e , and the conformational exchange contribution, R_{ex} , were obtained for this complex by Lipari-Szabo model-free analysis. The results show that VEK-30 displays distinctly different dynamic behavior as a consequence of binding to K2_{Pg}, manifest by decreased backbone flexibility, particularly at the binding region of the peptide. In contrast, the backbone dynamics parameters of K2_{Pg} displayed similar patterns in the free and bound forms, but, nonetheless, showed interesting differences. Based on our previous structure-function studies of this interaction, we also made comparisons of the VEK-30/K2_{Pg} dynamics results from different kringle modules complexed with small lysine analogs. The differences in dynamics observed for kringles with different ligands provide what we believe to be new insights into the interactions responsible for protein-ligand recognition and a better understanding of the differences in binding affinity and binding specificity of kringle domains with various ligands.

INTRODUCTION

The group A streptococcal surface virulence protein, PAM, a 43 kDa member of the M-like protein family, interacts with high affinity and specificity with the lysine binding site (LBS) of the 80-residue kringle-2 (K2) domain of human plasminogen (hPg) (1). A region of PAM, spanning amino acid residues 91–116, contains its $\alpha 1\alpha 2$ repeat sequences, and is responsible for binding of PAM to the K2_{Pg} domain, an event that allows for a proteolytic bacterial surface and heavily contributes to virulence of PAM+ group A streptococci (2). A functional internal peptide of PAM, namely VEK-30, derived from residues 85–113 of PAM, and containing the first and most of the second direct repeat, possesses a high-affinity binding site for K2_{Pg} ($K_D = 460$ nM). In contrast, there is no measurable affinity of VEK-30 to any of the other isolated hPg kringles (3–5).

Recently, we determined the NMR structures of the VEK-30/K2_{Pg} complex, which provided an understanding the intermolecular interactions that govern this interaction in solution (6). On the basis of the solution structure, as well as multiple sequence alignments of different kringle modules, and mutational studies, we proposed that the conserved hydrophobic binding core of K2_{Pg} (residues Tyr³⁵, Phe⁴⁰, Trp⁶⁰, Phe⁶², Trp⁷⁰, and Y⁷²), along with side chain residues in its anionic (Asp⁵⁴ and Asp⁵⁶), and cationic (Arg⁶⁹) centers, interacts with VEK-30 in a similar fashion as its binding to small lysine analogs, resulting in a basal level of binding affinity. However, several nonconserved residues, including Gly³⁴, Lys³⁹, Lys⁴³, and Arg⁵⁵, which specifically

exist in K2_{Pg}, and are located outside of the canonical LBS, strategically occupy specific positions that optimize interactions with VEK-30. These exosite interactions enhance affinity of VEK-30 to K2_{Pg} and facilitate the docking of VEK-30 to the K2_{Pg} domain by modulating ligand recognition and binding specificity.

Structural data alone are not sufficient for an in-depth prediction of binding properties (7–9). Internal dynamics commonly plays an important role in the function of proteins. For example, specific molecular recognition processes that occur in biological systems are dependent on the dynamic properties of the species involved. NMR spectroscopy is not only of great use for determining the atomic structures of proteins, but also is a powerful and unique tool for the study of dynamic properties of proteins at the atomic level (10–13). The dynamical analyses of relaxation data using the model-free approach introduce a global rotational correlation time (τ_m), an internal correlation time (τ_e), and an order parameter (S^2), the latter being related to the amplitudes of internal motions for each residue (14–16). These approaches are useful in furthering our understanding of the role of time-dependent conformational fluctuations involved in binding events, and provide insights into describing the contributions of motions on different timescales to the high affinity and binding specificity.

In this study, NMR relaxation data for both free and bound forms of VEK-30 and K2_{Pg} were collected and analyzed with Lipari-Szabo model-free approaches to investigate the changes in backbone dynamics induced by binding of each peptide. These dynamics studies provide important complementary information regarding the nature of this specific molecular recognition event.

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METHODS

Protein expression and purification

¹⁵N-K2_{Pg}[C4G/E56D/L72Y], a triple variant of wild-type (WT)-K2_{Pg} that displays enhanced affinity for lysine analogs and VEK-30 compared to WT-K2_{Pg} was expressed in *Pichia pastoris* GS115 cells and purified as described (6). Final yields of ¹⁵N-K2_{Pg} were typically 75–100 mg/L. ¹⁵N-VEK-30 was expressed in *Escherichia coli* and purified as published (6). Final recoveries of peptide were 10–15 mg/L.

The purity and correct folding of materials were verified by SDS-PAGE and 2D HSQC spectroscopy, respectively. MALDI-TOF mass spectrometry was used for molecular weight analyses. For the uniformly labeled ¹⁵N-peptides, single mass peaks were obtained at the correct molecular weights, indicating nearly complete incorporation of ¹⁵N.

NMR sample preparations

Samples (1 mM) were dissolved in a buffer containing 50 mM Hepes (pH 7.0), 1 mM EDTA, 200 mM NaCl, 3 mM NaN₃, 10% ²H₂O and 0.2 mM 2,2-dimethyl-2-silapentane-5-sulfonic acid. Samples of the binary complex were prepared at 1 mM (1:1 stoichiometry of VEK-30/K2_{Pg}).

NMR spectroscopy and backbone ¹⁵N relaxation measurements

NMR spectra were collected on a Bruker AVANCE 800 MHz spectrometer using a 5-mm triple resonance cryoprobe. The backbone chemical-shift assignments were obtained by a set of 3D-HNCA, HNCO, HNCACB, HBHA(CO)NH, and CBCA(CO)NH experiments (6). Data were zero-filled to double the original data points and apodized with 90° to 60° shift square sine bell window function before Fourier transformation. ¹H and ¹⁵N chemical shifts were referenced indirectly to internal 2,2-dimethyl-2-silapentane-5-sulfonic acid (17).

The backbone ¹⁵N relaxation parameters, including longitudinal relaxation rates (R_1), transverse relaxation rates (R_2), and steady-state heteronuclear [¹H]-¹⁵N NOEs, for free and complexed VEK-30 and K2_{Pg} were measured using standard pulse sequences at 298 K. All ¹⁵N- R_1 and ¹⁵N- R_2 relaxation experiments were carried out in an interleave manner with a 2-s recycle delay between scans. The relaxation delays used for complexed and free K2_{Pg} were 10 × 2, 100, 300, 500, 800, 1000, 1200, and 1600 ms for the R_1 experiments and were 17 × 2, 34, 51, 68, 85, 102, 119, 136, 153, 170, 204, and 238 ms for the R_2 experiments. For free VEK-30, relaxation delays were set to 10 × 2, 50, 90, 220, 350, 520, 680, 820, 960, and 1100 ms for the R_1 experiments and 17 × 2, 34, 51, 68, 85, 102, 119, 136, 153, 170, and 204 ms for the R_2 experiments. Duplicate spectra were used to estimate experimental errors. The relaxation rates were determined by fitting the crosspeak intensities to a single exponential function using nonlinear least-squares. The error in the rate constant was assessed from Monte Carlo simulations. [¹H]-¹⁵N NOE experiments were carried out in the absence and presence of a 3-s proton saturation period before the ¹⁵N excitation pulse, using recycle delays of 4 and 7 s. Heteronuclear NOE values were obtained from the ratios of the peak intensities measured with and without proton saturation. Peak intensities uncertainties were estimated from the noise level of the spectra (NOE measurements).

RESULTS

To investigate the motional properties of K2_{Pg} and VEK-30, as well as to obtain further insights into the molecular mechanisms of their mutual recognition, backbone ¹⁵N longitudinal relaxation rates (R_1), transverse relaxation rates (R_2), and heteronuclear ¹H-¹⁵N NOE values, for free and bound peptides, were measured.

¹⁵N relaxation measurements for K2_{Pg}

In these analyses, 66 and 69 of 87 residues were used for the free and the VEK-30 bound forms of K2_{Pg}, respectively. The unanalyzed residues included seven proline residues (Pro³⁰, Pro³⁷, Pro⁴¹, Pro⁵³, Pro⁵⁹, Pro⁶⁶, and Pro⁷⁶) that have no amide protons. Other residues were excluded due to weak signals or severe spectral overlap. The experimental relaxation rates, R_1 , R_2 , and ¹H-¹⁵N NOE values, are plotted against the amino acid sequence (Fig. S1 in the Supporting Material). Overall, the R_1 , R_2 , and NOE values of K2_{Pg}, complexed to VEK-30, show similar patterns in comparison to those of the apo form, i.e., there is little variation in R_1 value across the sequence of K2_{Pg}, and the R_2 rates exhibit clear differences between the bulk of protein and several regions. Further, a significant decrease in R_2 is observed at both the N- and C-termini. Relatively high NOE values appeared in the core region (1–78) with the large deviations from average values also observed at the two termini.

Average R_1 , R_2 , and NOE values of 1.34 s^{−1}, 11.9 s^{−1}, and 0.79, respectively, were obtained for free K2_{Pg}, and average R_1 , R_2 , and NOE values of 1.19 s^{−1}, 14.0 s^{−1}, and 0.80, respectively, were obtained for bound K2_{Pg}. The high backbone ¹H-¹⁵N NOE values indicate that K2_{Pg} is generally rigid on the ps timescale, both in its apo- and VEK-30-bound states. Residues in the two terminal segments of Tyr⁷–Glu¹ and Asp⁷⁴–Ala⁸⁰ showed relatively low R_2 and NOE values, indicating their high flexibility on the ps-ns timescale. These observations are consistent with the lack of observed medium- and long-range NOEs within these peptide regions, based on the NOE data (6). For both forms of K2_{Pg}, Gly⁶, Cys²², His³¹, Ala³², Lys³⁹, Thr⁶³, and Thr⁶⁴ show larger than the average R_2/R_1 values, indicative of conformational exchanges on μs-ms timescale and/or internal motions on the ps-ns timescale. Notably, the segment Gly³⁴–Lys⁴⁶ in the apo form manifests more fluctuations in R_2 rates than in the complexed state (Fig. S1).

¹⁵N relaxation measurements for VEK-30

Reliable relaxation parameters were obtained for 26 and 25 of 32 nonproline residues for apo and bound VEK-30, respectively. The unanalyzed residues were due to partial overlap or line-broadening, indicative of internal motion and/or rapid exchange with solvent. The residue-specific relaxation parameters are shown in Fig. S2. Overall, the internal dynamics of VEK-30 complexed to K2_{Pg} is significantly changed relative to apo-VEK-30. In the complex, the R_1 rates significantly decreased, and the R_2 rates systematically increased, compared with free VEK-30, in agreement with the almost three-fold increase in molecular mass of VEK-30 in the complex. The [¹H]-¹⁵N NOE value of free VEK-30 is systematically low (average value = 0.25 ± 0.34), compared to that of bound state (average value = 0.60 ± 0.33), suggesting that the functional peptide is highly mobile and

unstructured in absence of K2_{Pg}, which is consistent with the chemical shift index of VEK-30. The complex structure determined by x-ray and NMR methods showed that VEK-30 adopts an extended α -helix conformation throughout residues Ala⁶-Leu²⁶, whereas the N- and C-terminal regions are solvent exposed and structurally disordered. Most of the intermolecular interactions occur between a single face of the α -helix of VEK-30 (residues 6–21) and the LBS of K2_{Pg}. Consistent with the structure of VEK-30 in the complex, the bound peptide exhibits distinct differences in the relaxation parameters, R_2 and NOE in three regions. These include: N- and C-terminal residues, 1–5 and 27–30, respectively, which have lowest average R_2 ($5.5 \pm 2.1 \text{ s}^{-1}$); residues 6–21, which make the majority of close contacts with K2_{Pg}, and have the highest average R_2 value ($13.6 \pm 1.4 \text{ s}^{-1}$); and residues 22–27, which is an extended α -helical region, and has slightly lower R_2 values than residues 6–21. Similarly, the different regions of the bound peptide clearly show different ranges of NOE values. Both the N- and C-terminal residues have negative, or relatively low, NOE values as compared to the average NOE for the remainder of the peptide. Unlike the other relaxation measurements, the different regions of the peptide did not exhibit systematic variations in R_1 values. Therefore, Fig. S2 indicates that binding of K2_{Pg} perturbs the dynamics of VEK-30 by reducing backbone flexibility throughout the extended α -helix, more strikingly, at the main binding region of the residue 6–21 region, as demonstrated by changes in the overall patterns of the R_1 , R_2 , and NOE values plotted as a function of residue number.

Model-free analysis

The analysis of backbone dynamic parameters was carried out using the Lipari-Szabo model-free method, as implemented in the program, Tensor 2 (18). In this approach, relaxation rates were first used to estimate the diffusion tensor. Here, common procedures were followed to determine the rotational diffusion tensors of both apo-K2_{Pg} and VEK-30/K2_{Pg} by excluding residues with conformational exchanges and/or fast internal motions. Residues that satisfied these criteria were used to characterize the diffusion tensor, which was chosen over isotropic and anisotropic models. On the basis of the observation that the degree of anisotropy was found to be small ($D_{||}/D_{\perp} \approx 1.03$ and 1.06 for apo-K2_{Pg} and VEK-30-bound K2_{Pg}, respectively) and the improvement in χ^2 was not meaningful according to a statistical F-test, the diffusion tensors were best defined by isotropic rotational tumbling for both apo-K2_{Pg} and K2_{Pg} bound to VEK-30, thus providing the initial estimate of global correlation times (τ_m) of $6.72 \pm 0.02 \text{ ns}$ and $7.80 \pm 0.02 \text{ ns}$ for free K2_{Pg} and the K2_{Pg}/VEK-30 complex, respectively. The correlation times indicate that both forms of K2_{Pg} are in the monomeric state under the NMR experimental conditions, which is in accord with analytical ultracentrifugation data that we reported earlier (6).

The model-free parameters were extracted using the experimentally determined ¹⁵N relaxation rate parameters and the heteronuclear NOE values, and an isotropic diffusion model was used in the analysis. The amide bond length was fixed at 1.02 \AA , and a ¹⁵N chemical-shift anisotropy value of -175 ppm was used in the calculations. Five simplified models of internal mobility are defined. These are: model (M)1- S^2 ; model (M)2- S^2 , τ_e ; model (M)3- S^2 , R_{ex} ; model (M)4- S^2 , τ_e , R_{ex} ; model (M)5- S_f^2 , S_s^2 , τ_e . These models were iteratively used to fit the experimental data until the confidence reached 95% (19). Here, S^2 ($= S_f^2 S_s^2$) is the square of the generalized order parameter characterizing the amplitude of the internal motions, where S_f^2 and S_s^2 are the squares of the order parameters for the internal motions on the fast and slow timescales, respectively. τ_e is the effective correlation time, which indicates the timescale of internal motions. The confidence levels were estimated using 300 Monte Carlo simulations per run in combination with χ^2 and F-test criteria. To take into account the contribution to the experimental ¹⁵N R_2 relaxation rate from conformational exchange processes, an additional parameter (R_{ex}) was introduced. Using the model-free formalism, backbone amides were fitted to either one of the five models. Fig. S3 summarizes the number of residues that fit to each of the five models for both the apo- and VEK-30 bound K2_{Pg}. The optimized internal mobility parameters of the generalized order parameter, S^2 , the fast internal motion on ps-ns timescales, τ_e , and the conformational exchange, R_{ex} , on μ s-ms timescales are shown in Fig. 1 and Fig. 2.

Model-free analysis of backbone motions in K2_{Pg}

As shown in Fig. S3, after model selection, 34 residues for apo-K2_{Pg} were described by M1 and 10 residues were fit to M2, indicating internal motions (τ_e) on ps-ns timescales. A total of 11 residues were fitted to M3 and six residues were assigned to M4, suggesting more complicated and flexible internal motions. Five residues were assigned to M5. There are three residues that could not be fit to any of these models. Of the bound K2_{Pg} residues that were analyzed, 35 were fit to M1, 13 to M2, 11 to M3, 4 to M4, and 6 to M5. Two residues could not be fit to the five traditional models. Therefore, for both forms of K2_{Pg}, the majority of residues were best-fit with the simplest model, either with S^2 (M1) or S^2 and τ_e (M2). The extracted dynamic parameters (S^2 , τ_e , and R_{ex}) for free and bound forms of K2_{Pg}, are reported in Fig. 1. Average S^2 values of 0.88 ± 0.10 and 0.91 ± 0.09 were obtained for the free and VEK-30-bound states of K2_{Pg}, respectively, indicating that the K2_{Pg} domain, both in the absence and presence of VEK-30, shows similar restricted motions on ps-ns timescales (Fig. 3, A and B). For the core-region (residue 1–78), as compared in both states, $\Delta S^2_{\text{bound-free}}$ averages 0.028 . This suggests that VEK-30 binding results in a slight enhancement of overall rigidity on the ps-ns timescale. Despite this fact, several

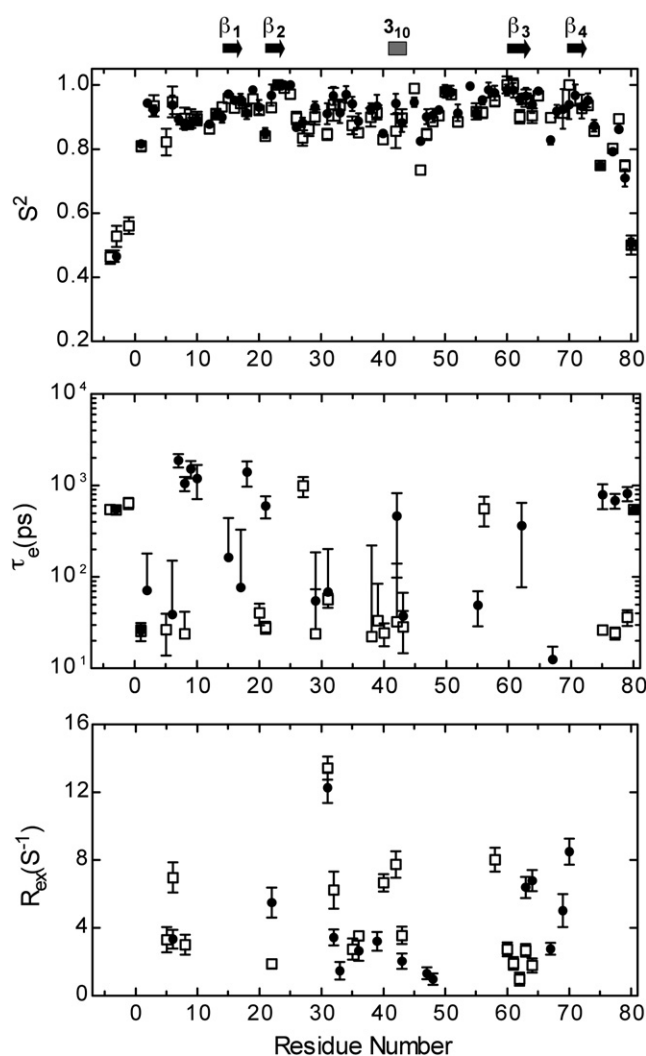


FIGURE 1 Residue-specific dynamics parameters calculated from the extended Lipari-Szabo model-free formalism for K2_{Pg} in absence (open squares) and presence (solid circles) of VEK-30. Secondary structure elements are displayed above the top graph.

residues located within, or in close proximity to, the binding center of apo-K2_{Pg}, e.g., Cys⁵⁰, Arg⁵¹, Arg⁵⁸, Trp⁶⁰, Cys⁶¹, Thr⁶³, Asp⁶⁵, and Trp⁷⁰, exhibit relatively high order parameters, suggesting that low mobility in these positions likely plays a role in restricting the conformational changes required for complex formation.

A total of 21 and 23 residues of apo- and VEK-30 complexed K2_{Pg}, respectively, undergo fast internal motions on the ps-ns timescale, as shown by τ_e values in the range of 20–1000 ps (free) and 30–1900 ps (complexed). Most of these residues have surface-exposed side chains. For some residues outside this binding surface, the mobility of K2_{Pg} on this timescale changes on binding to VEK-30. Notably, in the N-terminal electrostatic field region of Ser³⁸-Lys⁴³, more residues exhibit nonzero τ_e contributions for apo-K2_{Pg} (five residues are accounted for with models 2 or 4 in apo-form whereas only two residues require τ_e contributions

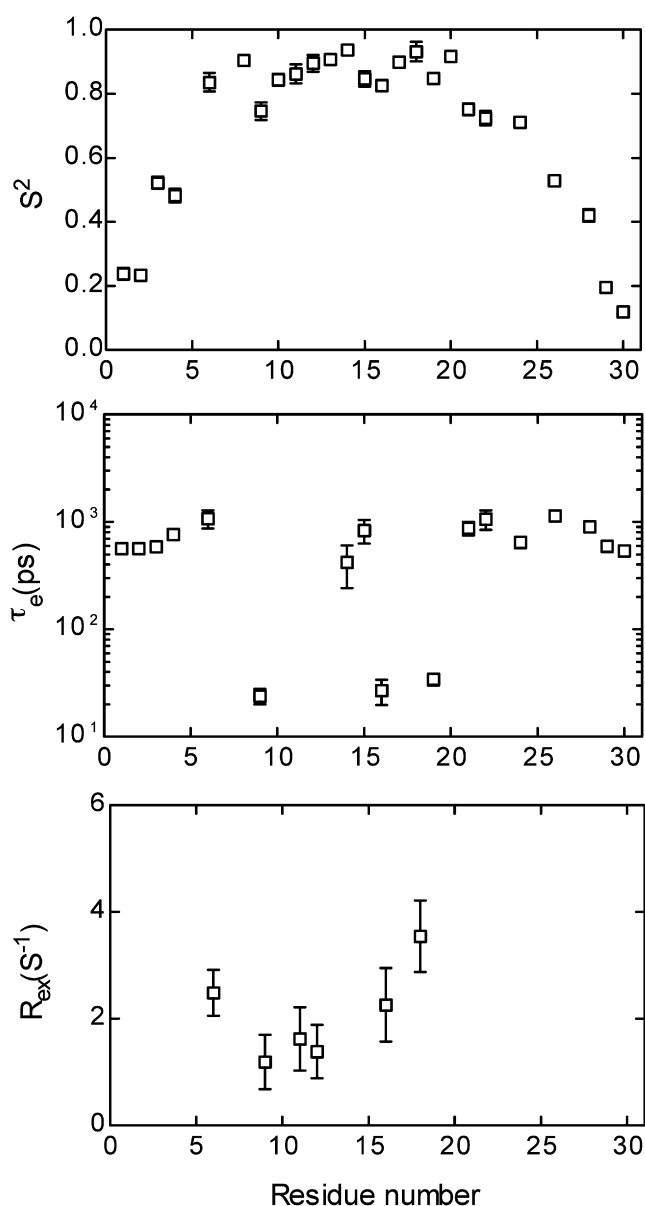


FIGURE 2 Residue-specific dynamics parameters calculated from the extended Lipari-Szabo Model-free formalism for VEK-30 binding to K2_{Pg}.

in bound form). This indicates that complexation of K2_{Pg} with VEK-30 reduced those internal motions characterized by correlation times of 20–460 ps in this region. In addition, slow motions, as reflected by R_{ex} in the μ s-ms timescale, are distributed throughout the backbone, especially around the binding sites of both forms (Fig. 3, C and D). In total, the R_{ex} parameter is required for 15 residues of bound K2_{Pg}, whereas 18 residues contain R_{ex} contributions in case of apo-K2_{Pg}. However, some differences were observed in several regions for conformational exchanges: 1), N-terminal residues Ser⁵-Asn⁸ showed conformational exchanges in apo-K2_{Pg}, whereas only Gly⁶ exhibited conformational exchange in the bound form; 2), a number of conformational exchange contributions were observed in the N-termini of

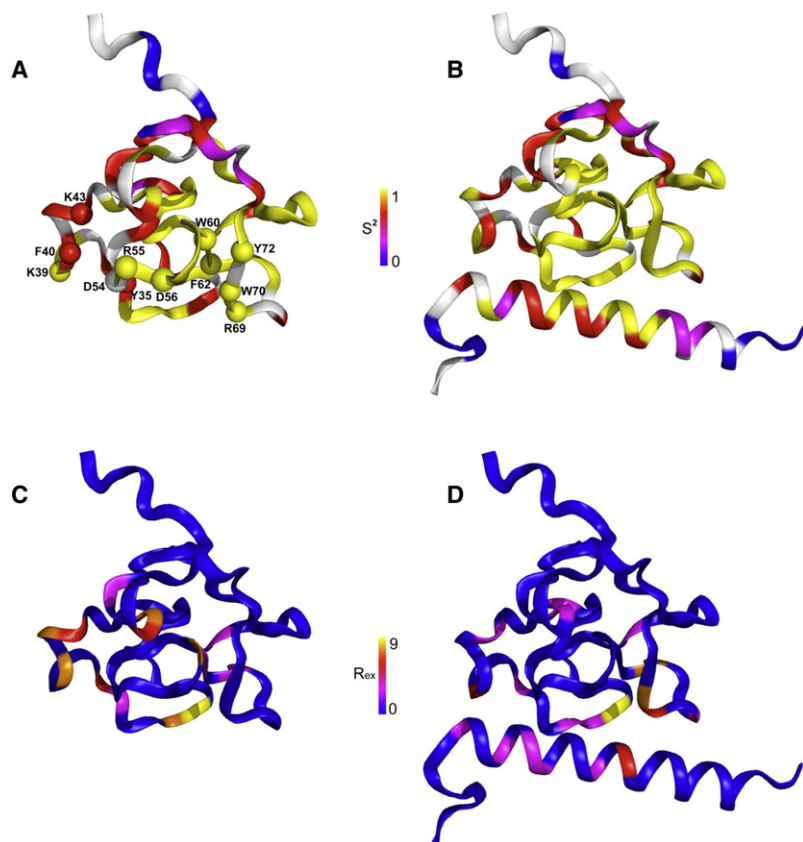


FIGURE 3 Mapping of the dynamics parameters on the protein structures both of free and bound forms with continuous-scale colors. (A and B). Ribbon diagrams of (A) apo-K2Pg and (B) VEK-30/K2Pg representing the generalized order parameter, S^2 , with color coding as follows: blue, $0 < S^2 < 0.7$; magenta, $0.7 < S^2 < 0.8$; red $0.8 < S^2 < 0.9$; yellow, $0.9 < S^2 < 1$. Unanalyzed residues are white. (C and D) Ribbon diagrams of apo-K2Pg (C) and VEK-30/K2Pg (D) representing the conformational exchange dynamics on the μ s-ms motional timescale, as indicated by R_{ex} (Hz). Color coding of R_{ex} is as follows: blue, $R_{ex} \sim 0$; magenta, $1 < R_{ex} < 3$; red, $3 < R_{ex} < 6$; orange, $6 < R_{ex} < 9$; yellow, $R_{ex} > 9$. Molecular models were drawn using the NMR average structure (PDB entry 2KJ4). The hydrophobic core, composed of Tyr³⁵, Phe⁴⁰, Trp⁶⁰, Phe⁶², Trp⁶⁷, and Phe⁷², and the anionic (Asp⁵⁴ and Asp⁵⁶) and cationic (Lys³⁹, Lys⁴³, Arg⁵⁵, and Arg⁶⁹) centers are shown in spheres with labeled residues (the color coding of the spheres is the same as the dynamics parameters) in Fig. 3 A.

the interacting interface between K2Pg and VEK-30 (Ser³⁴–Lys⁴³) for both forms, but the free form showed a higher average value ($4.8 \pm 2.2 \text{ s}^{-1}$) than the bound form ($2.3 \pm 0.7 \text{ s}^{-1}$) in this region; 3), in the absence of VEK-30, Arg⁵⁸, a residue near the anionic center, exhibited significant conformational exchange. Further, the critical hydrophobic binding region Trp⁶⁰–Phe⁶², also exhibited nonzero conformational exchange contributions. In contrast, these conformational exchanges substantially disappeared on binding to VEK-30, suggesting complexation reduces the low-frequency motions in the central binding region; and 4), VEK-30-bound K2Pg showed enhanced conformational exchange motions in the C-terminal binding region (Thr⁶³–Trp⁷⁰) compared to the free form. In particular, the binding sites at Arg⁶⁹ and Trp⁷⁰ had restricted internal motions in the free form, as reflected by the high S^2 value, whereas they displayed marked conformational exchange motions on binding to VEK-30, with R_{ex} of $5.0 \pm 0.9 \text{ s}^{-1}$ and $8.5 \pm 0.8 \text{ s}^{-1}$, respectively.

Model-free analysis of VEK-30 backbone motions in complex with K2Pg

Relaxation data for the VEK-30/K2Pg complex was analyzed assuming that the overall rotational diffusion is isotropic. The dynamics model best describing the data for each residue in bound VEK-30 is shown in Fig. 2. The S^2 values

for the N- and C-terminal residues are distinctly smaller than the S^2 values of other regions of the peptide. This indicates that both the N- and C-termini exhibit high mobility on the ps-ns timescale, which is consistent with relatively unstructured portions of the bound peptide. The average S^2 of the main binding region residues, 6–20, is 0.87 ± 0.05 , indicating limited ps-ns timescale motion. In contrast, the average S^2 value for extended region of α -helix, segment 21–26, is 0.68 ± 0.10 , which indicates that this region has increased mobility relative to the main binding region, perhaps due to the lack of contacts with K2Pg. In addition to the two termini of the peptide, several residues, viz., Ala⁶, Glu⁹, Lys¹⁴, Asn¹⁵, Glu¹⁶, Glu¹⁹, Ala²¹, Glu²², Glu²⁴, and Leu²⁶, show fast and slow internal motions (τ_c) that occurred on a 20 ps to 1.2 ns timescale, which are characterized by the complex models 2, 4, and 5, respectively.

Chemical shift perturbation on complexation

We have measured 2D ¹H-¹⁵N HSQC spectra of the peptide species involved in free and bound states (Fig. 4) to obtain direct insights into the effects of complex formation on protein structure due to the exquisite sensitivities of chemical shifts to the local environment. Fig. 5, A and B, shows the composite ¹H and ¹⁵N chemical shift changes versus residue numbers, and the mapping of the chemical shift changes onto the structure of complex (Fig. 5 C). Relatively slight

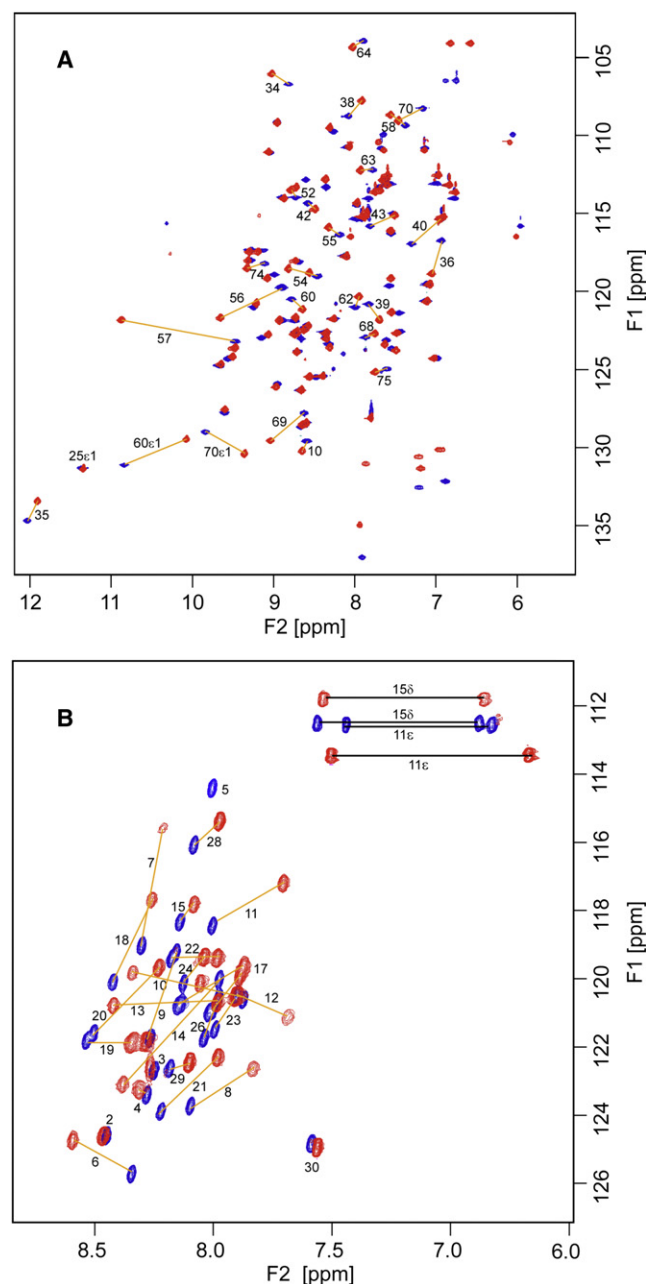


FIGURE 4 ¹H-¹⁵N HSQC spectral analyses of (A) K2_{Pg} and (B) VEK-30, respectively. Spectral shifts in the absence (red) and presence (blue) of VEK-30 (A) and in absence (blue) and presence (red) of K2_{Pg} are superimposed. The residue numbers are labeled in the plots.

perturbations were observed for most residues of K2_{Pg}, which are spatially far from the ligand-binding sites. In contrast, pronounced perturbations occurred in the vicinity of several binding regions (Fig. 5 C). The K2_{Pg} segments, Gly³⁴–Phe⁴⁰, Asp⁵⁴–Trp⁶⁰, and Arg⁶⁹–Trp⁷⁰, are the most sensitive to ligand binding, as revealed by the significantly perturbed chemical shifts (Fig. 5 A). They include residues located close to the hydrophobic (Gly³⁴, Tyr³⁵, Phe⁴⁰, Trp⁶⁰, Trp⁷⁰), cationic (Lys³⁹, Lys⁴³, Arg⁵⁵, R⁶⁹), and

anionic (Asp⁵⁴ and Asp⁵⁶) centers. Other residues exhibiting distinct binding-induced resonance shifts are Asp¹⁰, Ser¹⁴, Asp³³, Cys⁷³, and Ile⁷⁵, suggesting that residues far from the binding interface are affected by complexation. In addition, the side chain indole rings of Trp⁶⁰ and Trp⁷⁰ in the LBS also display chemical shift changes of aromatic NH^{ε1} resonances on binding VEK-30. Interestingly, a notable change was also observed for the side chain NH^ε resonance of Arg⁶⁹ (Fig. 4 A).

In case of VEK-30 binding to K2_{Pg}, almost all amide resonances of VEK-30 appeared in new positions in the HSQC spectra compared to uncomplexed VEK-30, except for the two termini (Val¹–Ser⁵ and Glu²⁹–Tyr³⁰) (Fig. 4 B). This shows that the majority of residues of VEK-30 display significant chemical-shift changes, demonstrating that their local environments were markedly affected on binding of K2_{Pg} (Fig. 5 C). The residues of the helical region are divided into two groups. One group includes residues, which participate directly in binding, exhibit >0.2 ppm in its chemical perturbations. The most striking differences occurred in residues Asp⁷, Glu⁹, Leu¹⁰, Leu¹³, Lys¹⁴, His¹⁸, and Glu²⁰. Another group of residues exhibiting distinct differences in amide chemical shifts includes Ala⁸, Gln¹¹, Arg¹², Asn¹⁵, Glu¹⁶, Glu¹⁹, and the segment, Glu²²–Lys²⁷. The side chains of Gln¹¹ and Asn¹⁵ also show significant differences in this regard (Fig. 4 B).

DISCUSSION

Whereas high-resolution structures of the complex of VEK-30/K2_{Pg} have been determined, no detailed studies of the dynamics of this interaction have been carried out. This has now been accomplished in the current study.

The backbone relaxation data on K2_{Pg}, and the subsequent model-free analysis, show that most amino acid residues in both its apo and bound forms have relatively high order parameters despite a paucity of regular secondary structure in K2_{Pg} (Figs. 1 and 3). This indicates that the backbone internal motions are highly restricted, and that K2_{Pg} is a rigid molecule in both forms. This is consistent with reports for related kringle modules (20–22). The rigid structure and restricted dynamical properties seem to be a general feature of kringle modules and confirm other reports that concluded that the LBS was preformed in kringle domains.

Effect of VEK-30 binding on the K2_{Pg}

Our results show that the backbone dynamics of K2_{Pg} are similar in the free and bound forms, but, nonetheless, interesting differences are present. We have analyzed these subtle differences in mobility based on known structural data, and conclude that by comparing the order parameters of free and bound K2_{Pg}, the binding of VEK-30 results in extensive increases and a lesser decreases in S², suggesting that binding of VEK-30 decreases the internal backbone motions

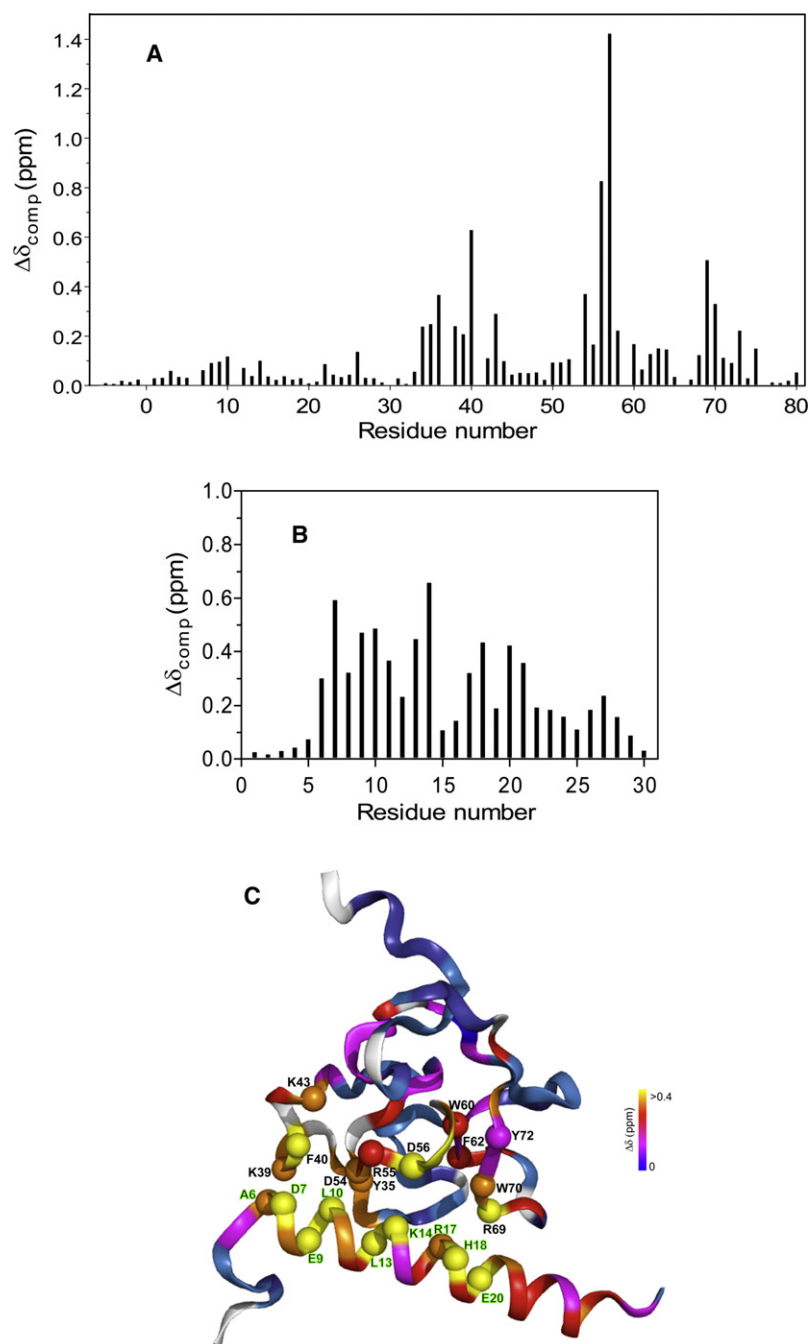


FIGURE 5 Chemical-shift differences between free and bound forms of K2_{Pg} (A) and VEK-30 (B) as a function of the residue number. Chemical-shift changes were calculated by the equation, $\Delta\delta_{\text{comp}} = [(\Delta\delta_{\text{HN}})^2 + (\Delta\delta_{\text{N}}/6)^2]^{1/2}$, where $\Delta\delta_{\text{H}}$ and $\Delta\delta_{\text{N}}$ represent the chemical-shift changes of ^1H and ^{15}N atoms between free and bound forms, respectively. (C) Color mapping of the chemical shift differences on the NMR structure (PDB 2KJ4): dark blue, $\Delta\delta_{\text{comp}} \sim 0$; light blue, $0.02 < \Delta\delta_{\text{comp}} < 0.06$; magenta, $0.06 < \Delta\delta_{\text{comp}} < 0.1$; red $0.1 < \Delta\delta_{\text{comp}} < 0.2$; orange, $0.2 < \Delta\delta_{\text{comp}} < 0.4$; yellow, $\Delta\delta_{\text{comp}} > 0.4$. Proline and unassigned residues are in white. The important binding sites in K2_{Pg} are shown in spheres as in Fig. 3 A. The residues of VEK-30 directly involved in binding are also shown in spheres with corresponding color coding and labeled by green font.

of K2_{Pg} (Fig. 6 and Fig. S4). A good example is noted in the region Tyr³⁵–Asn⁴², which experiences relative higher amplitude motions on the ps-ns timescale in the apo-form (average $S^2 = 0.87$). Binding of VEK-30 causes a systematic increase in the S^2 of this region ($S^2 = 0.91$), suggesting that more restricted motions occur herein on VEK-30 binding. We also conclude that slow motions on the μs -ms scale are biologically important because they are close to the timescales of functional processes, such as docking and protein folding (23–25). The comparison of the dynamic behavior of free and bound K2_{Pg} indicated that although high

frequency internal motions were similarly observed in most residues of K2_{Pg}, conformational exchanges in μs -ms timescales observed at the binding face of free and VEK-30-bound K2_{Pg}, have some different distributions and/or distinct tendencies. Notably, Phe⁴⁰, Asn⁴², Lys⁴³, Arg⁵⁸, and Trp⁶⁰ have high R_{ex} values in apo-K2_{Pg}. However, on binding of VEK-30, these prominent conformational dynamics on the μs -ms timescale are attenuated, and even quenched, suggesting that the conformational flexibility at these sites likely contributes to the recognition of the ligand and is required for binding. In contrast, the VEK-30-bound K2_{Pg}

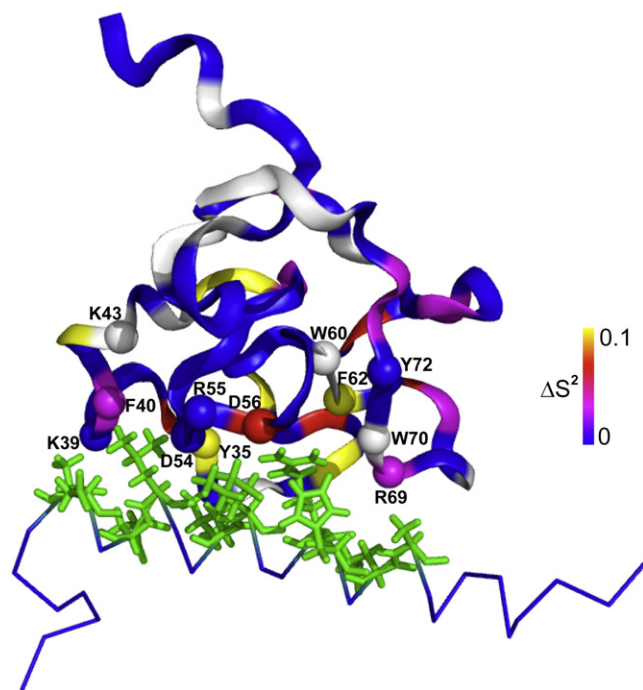


FIGURE 6 Effect of binding of VEK-30 on the generalized order parameter (S^2) of K2_{Pg}. $\Delta S^2 = S^2$ (after binding) – S^2 (before binding) and is mapped by continuous-scale color onto the structures. Positive ΔS^2 values denote enhanced rigidity of the protein backbone on binding. Residues with negative ΔS^2 values are white. The main chain of VEK-30 bound to K2_{Pg} is shown as a green ribbon and the side chains of residues involved in interactions with K2_{Pg} are represented as green sticks. Values of ΔS^2 are presented in Fig. S4. The important binding sites in K2_{Pg} are shown in spheres, as in Fig. 3 A, and the color coding of spheres is the same as ΔS^2 .

shows enhanced conformational exchange motions in the C-terminal binding region (Thr⁶³-Trp⁷⁰) compared to the free form. The increased mobility of residues on the μ s-ms timescale, as a result of ligand binding, has been observed in other complex systems (26,27). For this system, both the crystal structure reported previously (5) and the NMR solution structure (6) of VEK-30 bound to K2_{Pg} showed flexibility in the C-terminal binding region, especially two cationic residues, Lys⁶⁸ and Arg⁶⁹, which manifest different orientations with their long side chains, resulting in different binding modes with the three anionic residues (Glu¹⁶, Glu²⁰, and Glu²⁴) of VEK-30.

Comparison of the order parameters of free and bound K2_{Pg} showed that, unexpectedly, not only residues in the binding region of K2_{Pg}, but also some residues located spatially far from the binding site were affected by complex formation (Fig. 6). However, besides these changes, some of the dynamical features of the free form are retained in the complex. Therefore, the structure of apo-K2_{Pg} is comparatively rigid, but at the same time provides a relatively flexible interface for protein-ligand recognition. The flexibility at the binding interface is characteristic of the interaction surface of many proteins (28). The motions of residues in the vicinity of

the binding interface are constricted in part by the binding of VEK-30, indicative of stabilization their internal motions.

Effect of K2_{Pg} binding on dynamics properties of VEK-30

NMR spectral analysis showed that VEK-30 is unstructured and highly mobile in its apo form, whereas VEK-30 forms an extended regular α -helical structure when bound to K2_{Pg}. Here, we have shown that bound VEK-30 exhibits significantly different dynamical properties compared with free VEK-30. Interestingly, distinct backbone dynamical behavior is observed in different regions of the bound peptide, which correlate with specific interactions with K2_{Pg}. The residues at the two termini of VEK-30 have low S^2 values, whereas the majority of the helical residues in VEK-30 have relatively high order parameters. Mostly striking, the main binding region (residues 6–21) of VEK-30 displays the most restricted motion, reflected as the highest S^2 values. Analysis of the tertiary structure of VEK-30/K2_{Pg} show many hydrophobic and electrostatic interactions in addition to the intermolecular hydrogen bonding, which may restrict the internal motions of this region. In contrast, the extended part of helix (residues 22–27) has relatively low order parameters and exhibits increased dynamics relative to the main binding portion of the helix (residues 6–21) of this peptide. This suggests that this region of the peptide experiences higher amplitude motions on a ps-ns timescale. This is consistent with the structure of the complex, which indicates little or no interactions of the C-terminal portion of VEK-30 with K2_{Pg}. Model-free analyses show that among the binding region residues (residues 6–21) of VEK-30, five residues are best fit by model 1, suggesting that this binding region exhibits a much simpler motion and is more immobilized on binding. At the same time, six nonzero R_{ex} and numerous τ_e contributions appeared, indicating that flexibility of the conformation of VEK-30 may be required for high-affinity and high-specificity of the binding. In summary, binding of VEK-30 to K2_{Pg} results in widespread rigidification of the backbone in the extended helix (Ala⁶-Lys²⁷) of VEK-30, despite the complex dynamics that exist in this region.

Comparison of the dynamics of other ligand/kringle interactions

The structures of kringle modules and their complexes with lysine analogs have been solved by both NMR and x-ray crystallographic methods (20,22,29–32). These structures reveal similar binding sites, involving hydrophobic and electrostatic interactions. Furthermore, dynamics studies show that they have similar patterns of internal motions on binding of lysine analogs (20,21,32). NMR spectra also indicated that almost no visible shifts are observed in kringle modules after binding of small lysine analogs, except for several

important binding sites (20,21). However, in contrast, our results show that more amide resonances of K2_{pg} in the HSQC spectrum appeared in new positions after binding to VEK-30. This indicates that K2_{pg} undergoes larger and more extensive perturbations when bound to VEK-30 compared to kringles bound to small lysine analogs, and likely involve exosite interactions distal from the LBS.

In addition to primary sequence alignments and structural comparisons, investigation of the dynamical behavior of different kringle domains, complexed with different ligands, contributes to explanations of the differences in binding affinities and binding specificities with these ligands, as well as the very tight and selective binding of VEK-30 to K2_{pg}. Despite the similar dynamical patterns of kringles in some regions, K2_{pg} exhibited somewhat different dynamics at multiple timescales, which likely contribute to the differences in its binding specificity. Distinct from other kringle modules, which display nearly no change in the S^2 for apo- and lysine analog-bound states, the S^2 for K2_{pg} bound to VEK-30 is slightly increased compared with the apo-form, indicating that the binding of VEK-30 rigidifies K2_{pg} to some extent. Another major difference is seen for some residues near the binding site of kringle modules, especially at exosites of the LBS that were confirmed to be involved in interactions with VEK-30, and also contribute to the stability of the VEK-30/K2_{pg} according to our solution structural analysis and mutagenesis studies reported earlier (6). Backbone dynamics of apo-K1_{pg} reported previously did not show significant chemical exchanges at exosites of the LBS (21). However, our present results show that the exosite residues of LBS in K2_{pg}, Tyr³⁵-Ile³⁶, Phe⁴⁰, Asn⁴²-Lys⁴³, Arg⁵⁸, and Cys⁶¹-Thr⁶⁴, experience chemical exchanges with R_{ex} values of 1.0–8.0 s⁻¹. These differences with other kringle modules suggest that the newly identified exosites of K2_{pg} adopt more flexible conformations in the absence of ligand, thus allowing local structural rearrangements to accommodate the binding of larger- and longer-chain peptides, an important feature in explaining the high specificity of K2_{pg} binding to VEK-30.

We reported recently the solution structure of the VEK-30/K2_{pg} complex, and compared the structures of K2_{pg} bound to the small ligand, t-aminomethyl-cyclohexane-1-carboxylic acid (AMCHA) (6). The results showed that although the NMR structure of K2_{pg} in the VEK-30/K2_{pg} complex is similar to that of K2_{pg} bound to AMCHA, some subtle differences were observed in side chain geometries of several binding site residues. Most strikingly, two cationic residues, Lys³⁹ and Lys⁴³, have distinctly different side chain orientations in the two complexes, and the topology of backbone in vicinity of Lys⁴³ is distinctly different. Comparing the dynamics of the two ligand-bound systems, restricted motions on ps-ns timescales are fitted for residues Cys²², His³¹, His³³, Ile³⁶, Thr⁶³, Thr⁶⁴, and Trp⁷⁰ of K2_{pg} with both ligands. However, consistent with their solution structures, the backbone dynamics also revealed some subtle

differences. It is interesting to note that these residues have nonzero R_{ex} values in VEK-30/K2_{pg} (Lys³⁹ and Lys⁴³ show a value of 3.2 and 2.1 s⁻¹, respectively), but not in the AMCHA/K2_{pg} complex, indicating the VEK-30 bound K2_{pg} displays more complex motions in these regions. In addition, comparisons of ligand-bound K2_{pg} structures show that the aromatic ring of Phe⁴⁰ in VEK-30/K2_{pg} is a somewhat closer to the ligand center, such that it makes hydrophobic contacts with the methyl groups of Leu¹⁰ and Leu¹³ of VEK-30. By investigating the dynamical behavior of Phe⁴⁰ in two complexed states, it is seen that this residue experiences flexible motions at the ps timescale, characterized by a relatively low order parameter (S^2 of Phe⁴⁰ is 0.77, whereas the overall generalized S^2 of K2_{pg} is 0.85), when complexed to AMCHA. In contrast, this residue becomes considerably more rigid on binding of VEK-30, displaying a higher S^2 value of 0.85. This is consistent with the fact Phe⁴⁰ directly participates in this binding event in the complex of VEK-30/K2_{pg}, but not in the complex of AMCHA/K2_{pg} (6).

An additional significant difference is observed in the C-terminal binding region of K2_{pg}. In the AMCHA/K2_{pg} complex, only Trp⁷⁰ exhibits conformational exchange on μ s-ms timescale, whereas three residues, specifically, Asn⁶⁷, Arg⁶⁹, and Trp⁷⁰, have significant conformational exchange rates when bound to VEK-30. There are some structural differences in the orientations of side chains of binding site residues, Lys⁶⁸ and Arg⁶⁹, due to the different interactions involved. This impacts the mobility of binding sites and vicinal residues. These differences are also reflected in HSQC spectra of K2_{pg} by comparison of chemical shift perturbations derived from binding of different ligands. Because the K2_{pg} used in these two complexes has a single site mutation at position 72 (Tyr⁷² and Leu⁷² are present in VEK-30/K2_{pg} and AMCHA/K2_{pg}, respectively), the above differences in the region of Asn⁶⁷-Trp⁷⁰ could be due to the effects of this mutation. In this regard, the EACA/K1_{pg}, AMCHA/K1_{pg}, and EACA/KIV₈ complexes were investigated regarding relaxation data in this region, which have the same residue, Tyr⁷², and are bound to the same or similar lysine analogs (21,22,32). However, the residues in this region did not exhibit chemical exchanges of high magnitude for any of these kringle/ligand systems. Therefore, the possible influence of the Leu⁷²Tyr mutation is excluded in the VEK-30/K2_{pg} complex and these substantial differences of backbone dynamics in this region are likely attributable to the different docking modes between lysine analogs and VEK-30.

The dynamical behavior of each component of the VEK-30 and K2_{pg} in their apo- and bound forms, and the dynamical comparisons with different kringle-ligand systems, together with our previous structural and mutagenesis studies, enable us to further probe the correlation between the dynamics and binding of VEK-30 bound to K2_{pg}. As shown in Fig. 5 C, the major binding interactions in

VEK-30/K2_{Pg} occur between each face of VEK-30 and K2_{Pg}. First, an exposed hydrophobic groove, consisting of Tyr³⁵, Phe⁴⁰, Trp⁶⁰, Tyr⁶², Trp⁷⁰, and Tyr⁷², extends throughout the binding interface, and correspondingly, several hydrophobic side chains of VEK-30, Ala⁶, Leu¹⁰, Leu¹³, Arg¹⁷, and Ala²¹, are buried within this groove, resulting in a broad hydrophobic interface between K2_{Pg} and VEK-30. Additionally, residues Asp⁵⁴ and Asp⁵⁶ of K2_{Pg} form an anionic locus interacting with Lys¹⁴, Arg¹⁷, and His¹⁸ of VEK-30. Also, Arg⁶⁹ of K2_{Pg} acts as a C-terminal cationic locus interacting with Glu¹⁶ and Glu²⁰ of VEK-30. The above extensive hydrophobic and ionic interactions were verified to be important to stabilization of the complex through site-specific mutagenesis studies (5), thus providing the molecular basis of the high affinity. These dynamical analyses show that, for K2_{Pg}, most of these binding sites and their vicinal residues undergo comparatively restricted motions in the ps-ns timescale in both apo- and its VEK-30 bound form. This indicates that the broad hydrophobic groove and the central and C-terminal electrostatic fields are both preformed and sufficiently rigid, to allow the K2_{Pg} binding site to accommodate the VEK-30 peptide. Interestingly, some residues near Arg⁶⁹ exhibit significant chemical exchange processes on binding of VEK-30, thus providing possibilities to interact with the carboxyl moiety of Glu residues of VEK-30 through local conformational adjustments in this region. Additionally, in the N-terminus of the interface, the exosites of the LBS, Lys³⁹, Lys⁴³, and Arg⁵⁵ of K2_{Pg}, engage the N-terminus of VEK-30 by electrostatic and hydrogen bond interactions with negatively charged residues, Asp⁷ and Glu⁹, of VEK-30, forming a sandwich electrostatic field. Our previous site-specific mutagenesis studies confirmed that this electrostatic field plays an additional important role in the binding of VEK-30 to K2_{Pg} (6). However, these dynamical studies show that the N-terminal cationic locus and its vicinal residues in apo-K2_{Pg} experience relative flexible internal motions on the ps-ns timescale and even higher amplitude chemical exchange processes on the μ s-ms timescale, and these motions are stabilized after VEK-30 docks to K2_{Pg}. That this phenomena did not occur in a related region of other kringle modules, suggests that these dynamical processes in the N-terminal interface play a role in the specific recognition of VEK-30 by K2_{Pg}.

In summary, whereas K2_{Pg} is relatively rigid with a preformed LBS, this allows only a basal level of binding to small molecules. A larger ligand, e.g., VEK-30, is a highly mobile and unstructured peptide in the absence of K2_{Pg}, suggesting that the binding of VEK-30 to K2_{Pg} results in a decreased entropy for both the VEK-30 and K2_{Pg}. Although it would be energetically unfavorable, presumably the flexibility in the unbound form allows the protein to maximize the binding enthalpy at the extended interface, overcoming the unfavorable entropic contribution. The flexibility of the peptide ligand and some of the exosite residues of K2_{Pg} allow for additional binding energy as compared to small

molecule ligands, thus explaining on a dynamic level the tight and specific binding of VEK-30 to human K2_{Pg}.

SUPPORTING MATERIAL

Four figures are available at [http://www.biophysj.org/biophysj/supplemental/S0006-3495\(10\)00484-4](http://www.biophysj.org/biophysj/supplemental/S0006-3495(10)00484-4).

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